FALL 2026

MOEBIUS SYNDROME FOUNDATION NEWS

WWW.MOEBIUSSYNDROME.ORG

"I want to exist in a world where my voice is a sword that fights on behalf of children who deserve more than what they have." – Heaven Baylor

A Season of AWARENESS, ADVOCACY, AND CONNECTION

Don't miss Heaven Baylor's "Trial by Fire: Moebius Syndrome, Social Work, & Law School" on page 4, part of our ongoing efforts to raise awareness and support advocacy efforts happening throughout our community.

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OUR MISSION

The mission of the Moebius Syndrome Foundation is to provide information and support to individuals with Moebius syndrome and their families, promote greater awareness and understanding of Moebius syndrome, and to advocate for scientific research to advance the diagnosis and treatment of Moebius syndrome and its associated conditions.

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MOEBIUS SYNDROME NEWS

newsletter@moebiussyndrome.org

The Moebius Syndrome News is published two times per year. Articles, photographs, questions and comments are welcomed. The Moebius Syndrome News reserves the right to edit submitted material. Opinions expressed in the Moebius Syndrome News are each author's and do not reflect that of the Moebius Syndrome Foundation or the Moebius Syndrome News.

The newsletter is on the website at: www.moebiussyndrome.org

greetings, MOEBIUS COMMUNITY

Dear Friends,

For many families, the hardest part of a Moebius syndrome diagnosis is not the medical information — it is the feeling of being the only one.

That is why we are glad to open this issue with Moebius Connections. The Foundation has been connecting people since our earliest days in 1994, and our more formalized peer program has quietly made introductions within this community since 2022. Over the past year, we rebuilt it with a clearer process, more thoughtful matching, and volunteers ready to walk alongside you. It is open now to anyone seeking a friend or mentor who truly understands.

The pages that follow show that same commitment at work in different forms: awareness efforts that change how the world sees us, scholarships that open doors, research that moves our understanding forward, fundraisers and volunteers who give generously of their time, and donors whose support makes all of it possible.

Whatever brought you to this community — your own diagnosis, your child's, or a friendship that led you here — you belong. Thank you for being part of it.

With gratitude,
Rebecca Maher, President

YOUR BOARD OF DIRECTORS AT WORK: A 2026 SNAPSHOT

Everything MSF offers this community is shaped by a volunteer board that gives its time all year long. Here is what that work has looked like in 2026.

Board meetings. The board meets by video four times a year and gathers in person every year. This October, the full board travels to Pittsburgh for a weekend of strategic planning focused on how MSF can best support the Moebius syndrome community.

2027 conference. Board members are deeply involved in planning next year's national conference in Pittsburgh, from the largest decisions down to the smallest details.

Financial stewardship. The board approved the 2026 budget in January. The Finance Committee reviews financial reports monthly, with a full board review each quarter. In March, MSF engaged an independent auditing firm to complete a full audit of fiscal year 2025. The audit returned an unmodified opinion, the strongest result available, confirming that our financial statements are fairly presented and free of material misstatement. The board reviewed the findings and approved both the audit and the Form 990 filing.

Membership. In August, MSF launched Moebius Connect, a free app that gives members easier access to resources and to one another. The board also guided the evolution of our mentorship and friendship program, relaunched in September as Moebius Connections and coordinated by board members.

Philanthropy. Every board member is expected to support MSF's fundraising. This year's campaigns included Making A Difference for Moebius each April and Moves for Moebius in September and October. Board member Michael Mendenhall also organized Mulligans for Moebius, a charity golf tournament in Las Vegas that raised more than $40,000. Additional initiatives are planned before year's end.

Research. The board continues to support the Moebius Syndrome Foundation Research Registry, launched in 2025, and led this year's effort to update the definition of Moebius syndrome on the National Organization for Rare Disorders (NORD) website – www.rarediseases.org. NORD has since asked a board member and a Scientific Advisory Board member to work with OpenEvidence AI so that families searching for answers find accurate ones. Planning begins soon for the 2027 Scientific Research Symposium and CME event.

You can learn more about individual board members at https://moebiussyndrome.org/leadership/.

RAISING *awareness*

If you have Moebius syndrome, or you love someone who does, you already know how much a single interaction can shape a day. You have seen what it looks like when someone is dismissed, overlooked, or made to feel less than. You have also seen what changes in a person when they feel seen, respected, and included.

That shared experience is one of the things that connects our community. The other is what we do with it. We know more change is needed, and together we are working to make it happen.

We would like to highlight some of our community members who have worked to raise awareness about Moebius syndrome in the last few months. This isn't everyone, as we know there are many people raising awareness in many places and in many ways. Thank you to each of you!

David Freedman wrote an article called The Importance of Being Assertive, which speaks to the importance of learning to be assertive and realizing he didn't need every answer up front. He only needed to recognize that "I was enough," and trust that the tools for speaking up would come with time. You can read David's article and blag at https://freedmanonthego.com/

Dallas Lewallen raised awareness about Moebius syndrome at the annual Wagon Days Parade in Ketchum, ID during Labor Day weekend. Dallas and family walked down Sun Valley Road near all the fun prior to the parade and spoke with many people about Moebius syndrome.

Rhyan Nelson advocated for her ability to take a driver's course, adapted driving controls, and now has her license. Rhyan is a great example of overcoming obstacles when told she couldn't do something.

Moni Williams- Moni Williams received a certificate from the Detroit Mayor for raising awareness about Moebius syndrome. Congratulations to Moni!

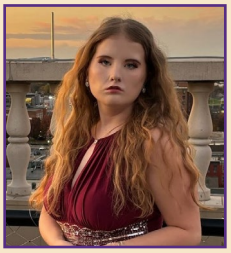
COMMUNITY EMPOWERMENT GRANT- OPEN THROUGH OCTOBER 30, 2026!

If you have a great idea that would help us fulfill our mission by expanding advocacy, awareness, or resources that would benefit the Moebius syndrome community, please apply. One or more people are awarded up to $2,000 each year. Learn more at https://moebiussyndrome.org/communitygrant/

This album is rooted in empowerment. In dancing through the pain, transforming challenges into strength, and finding joy in the places where it's often overlooked. With your support, this project has the power to reach people who need to feel seen, understood, and uplifted.

~Austin Halls, 2025 Community Empowerment Grant recipient

Austin's album, DISIBILI-TEA is available for streaming on Spotify and can be purchased and downloaded from Amazon.com.



Shelby Welch- July is Disability Pride Month, and Shelby Welch marked it by sharing a personal blog about what living with a disability has felt like for her this summer. She reflects on a semester where illness, daytime sleepiness, and communication barriers left her running on empty, and on what it took to finally request new accommodations. She writes candidly about being her own harshest critic: the instinct to push through, the belief that asking for grace means failing, and the discovery that disability burnout is real. She does not wrap it up with a neat answer. There is no single way to be disabled, and she is still learning what it looks like for her.

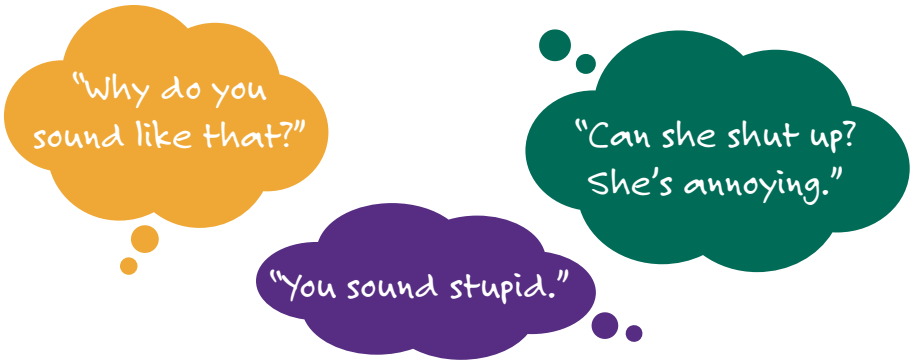
"There is no guide on how to be disabled because it means something different to everyone in the disability community."

READ SHELBY'S BLOG BY SCANNING THE QR CODE:

https://shelsjourney.wixsite.com/makeupdiamond/post/how-to-be-disabled

TRIAL BY FIRE: MOEBIUS SYNDROME, SOCIAL WORK, & LAW SCHOOL

By Heaven Baylor



Standing in front of three mock judges for 1L moot court at the University of Texas School of Law, these words, and the feelings they evoked, were unavoidable, even as I stared down at my oral argument outline. Consolidating my anxiety into a tidy box is an ongoing battle. While many people fear public speaking generally, I am deeply familiar with the shock and confused stares of people who expected "normal" and instead spoke to me, someone with Moebius syndrome, a neurological condition that impacts the cranial nerves. In my case, my tongue and bottom lip are the primary areas affected. My tongue is mostly paralyzed. I battle my anxiety, forged through years of unkindness, bullying, and fighting for my place in spaces people thought I didn't belong.

I know, as simply a matter of fact, that I sound different. However, when I talk, I hear a normal voice. Cameras and recordings expose the truth of the matter. Ever since I first heard my voice recorded, I've been keenly aware of it. I know that my mouth moves differently than yours, and my face, though not as obviously paralyzed as some of my community members', looks a bit different in its own small ways. My body can't accomplish all the same things a "normal" body can accomplish.

I know what it feels like to be
underestimated,
and I went through many phases growing up
where I had to fight to
make sense of myself
and my place in the world.

My journey to law school and my desire to advocate for other people were forged in metaphorical fire. I know what it feels like to be underestimated, and I went through many phases growing up where I had to fight to make sense of myself and my place in the world. Law school, so far, has been a similar experience. The irony of my desire to advocate for others is that my voice is one of my greatest insecurities when I'm around people I don't know well. As I've gotten older, I've spent lots of time trying to accept myself for what I am. Over time, I've gotten really good at handling other people's thoughts and expectations of me too. As a social worker, you hear a lot of stuff from a lot of people. And some people have no filter. That experience was invaluable to me before starting law school, though, and studying and practicing social work taught me how to communicate in these situations.

That being said, doing the actual work of becoming an advocate is terrifying. I hear the voices that have built up over the years, the self-doubt, the imposter syndrome, and the deep-seated feeling that I ended up somewhere I don't belong. I'm not the best public speaker, but I want to be better. I want to exist in a world where my voice is a sword that fights on behalf of children who deserve more than what they have. But along the way, I have to remind myself that it isn't my job to be someone else: a person who is more confident, more "normal," or more intelligent. I am who I am, and the only way I will get better at the things I am bad at is if I force myself to be uncomfortable. Law school is a trial by fire. It's so challenging partly because a room full of people who are used to being towards the tops of their classes are suddenly competing against each other for everything, and someone has to lose. It's the nature of the game due to the curve, and all activities have limited slots. If you've put stock in your academic abilities as part of your identity, what happens when suddenly you're in an environment where you can't rely on that anymore?

The work of becoming a lawyer has been enjoyable. It's almost been emotionally grueling, especially considering the fact that life has thrown its own curveballs that many of my classmates don't have to deal with. I've also realized that I don't know myself as well as I thought I did. I'm not sure what exactly I want for myself. More importantly, I'm starting to wonder where I fit in and whether my goals are still the same.

But either way, I will keep pushing through. I know that the Moebius community can do hard things. Growth happens in discomfort, and I've gotten better at learning how to sit with those feelings. In the spring, I will take trial advocacy, an entire class focused on 1.) improving your presence as an advocate and 2.) teaching trial skills. It is the one thing I know I really need to do, but I am scared to do it. Being judged based on my greatest insecurity sounds deeply uncomfortable.

That being said, if any young folks with Moebius are considering professional careers of any kind, I encourage you to take the leap. It can be unsettling at times, but we need more representation in these fields, and no one understands clients better than those who have shared similar lived experiences.

UPCOMING *events*

For more detailed information about MSF events, please scan the QR code to visit our events page.

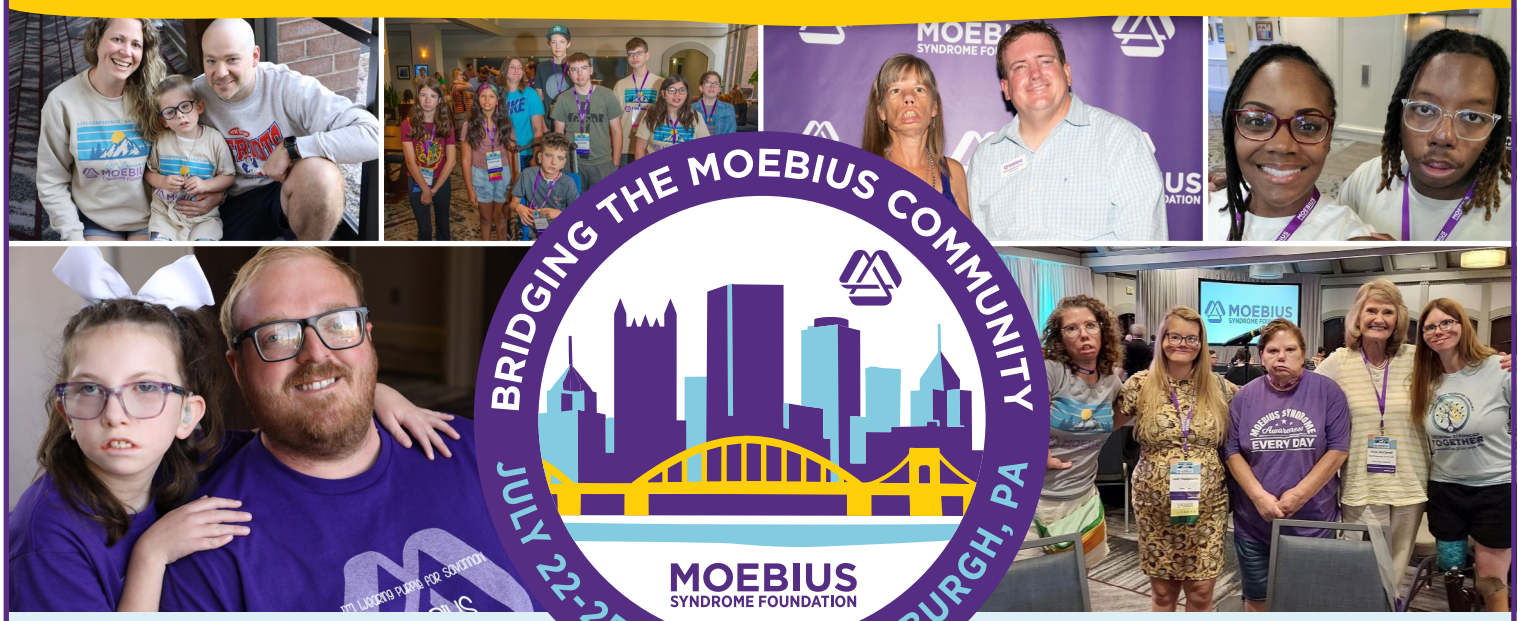
NOVEMBER 5 & 19	Grief Support Group
NOVEMBER 7	NC Kingfest and Group Dinner Regional Event
NOVEMBER 15	Pumpkins with Purpose – Kentucky & Tennessee Regional Event
DECEMBER 1	Giving Tuesday
DECEMBER 6	Just for Adults with Moebius Syndrome Chat
JANUARY 24	Moebius Syndrome Awareness Day (MSAD)

Keep a lookout for updates about MSAD events in January!

Our virtual gatherings (Just for Adults, Just for Adults: Midlife & Beyond, Spanish, LGBTQIA+ and Grief Support Group) will continue on a regular basis in 2027.

CALL FOR SPEAKERS

Moebius Syndrome Foundation Conference



Deadline to apply is November 6, 2026

Contact conference@moebiussyndrome.org with any questions.

BACK TO *school*

Students in the Moebius syndrome community are back in the classrooms and accomplishing great things. We are pleased to highlight outstanding students who received Moebius Syndrome Foundation Educational Scholarships this year, as well as a few students of all ages who are ready for the school year!

This last May, we were proud to recognize eight outstanding students who have worked hard to advance their education and pursue their goals. Each was awarded a $3,000 scholarship to support the 2026 fall semester's tuition and fees.

BACK-TO-SCHOOL

This August and September, students from all walks of life headed back to the classroom. The MSF received many requests for information and resources from parents sending their children back to school. Website resources are available year-round that can be shared with the entire school faculty. You can learn more about these resources by visiting our website at https://moebiussyndrome.org/school-resources-2/

We wanted to share these photos of three students heading to Pre-K, Middle School, and last year of college! We wish everyone a great school year.

RESOURCES TO SUPPORT THE MOEBIUS SYNDROME COMMUNITY

Learn more about these and other resources at moebiussyndrome.org.

ADVOCACY & AWARENESS

- Employment Resources (available now)
- Moebius Connect App (download for free at Apple Store and Google Play Store)
- Newsletters (sign-up now)
- Research updates and information (learn more now)
- Research Grants (also financial resources)
- School Resources (available now)

FINANCIAL RESOURCES

- Community Empowerment Grant (apply now - October 31, 2026)
- Conference Scholarships (opens next in early February 2027)
- Educational Scholarships (opens next in January 2027)
- Free & Low Cost Dental Resources (available now)
- Research Grants (opens next in 2027)
- Travel Reimbursement (apply now)

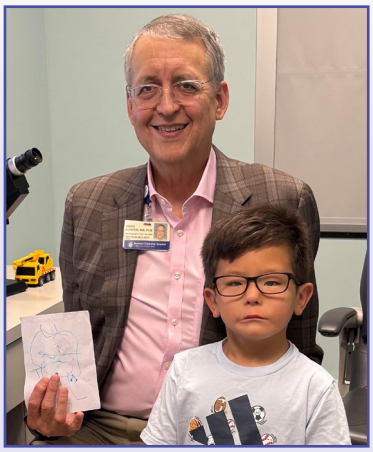
EDUCATION

- Materials to share information with Healthcare Providers and Educators
- New Member Packets (available to new parents of a child with Moebius, and adults with Moebius now)
- Research Updates and Articles
- Webinars and virtual events

CONNECTION

- Conferences (next national conference will be in Pittsburgh, PA on July 22-25, 2027)
- Events (check out upcoming events now)
- Moebius Connections (sign-up today)
- Social Media Support Groups (join our private Facebook group: The Home for the Moebius Community now)

Please contact us if you have any on how to access any of the resources listed above.
info@moebiussyndrome.org or call (844) 663-2487 ext. 704



Dr. David Hunter and Jackson Roberts
Photo credit: Patricia Roberts

THE ROBERTS FAMILY SHARES THEIR EXPERIENCE WITH THE MEDICAL TRAVEL REIMBURSEMENT PROGRAM

Jackson is five years old living with Moebius syndrome in Bucks County, Pennsylvania. When he was just two years old, we started traveling to Boston Children's Hospital to meet with Dr. Hunter. We have traveled to Boston once or twice a year for surgery, pre-and postop visits, along with regular routine eye exams.

The Moebius Syndrome Foundation reimbursement program took the financial burden off our hands this trip. We are forever grateful. Thank you so much to the Moebius Syndrome Foundation! Our hearts are full. Jackson's appointment went well, and we will continue to offer him every opportunity. We are proud to be involved with such an incredible organization and the Moebius community!

The Roberts Family

research UPDATES

MOEBIUS SYNDROME FOUNDATION TO SUPPORT THE OPENEVIDENCE AI PROJECT

Families often find themselves explaining Moebius syndrome to the very clinicians treating it. That is why it matters where doctors go looking for answers.

The National Organization for Rare Disorders (NORD) has been helping families and physicians find information about rare disorders for over 40 years. Their first textbook, "A Physician's Guide to Rare Diseases," was transformed into an online Rare Disease Database and has been one of the most widely used references of its kind. This database will once again be updated.

On March 12, 2026, the National Organization for Rare Disorders (NORD) officially partnered with OpenEvidence AI. Often described as a "ChatGPT for doctors" but with rigorous citations, OpenEvidence AI synthesizes peer-reviewed medical literature reviews in response to inquiries about rare diseases.

NORD has invited Kathy Kraus, MD, MSF Board Member, and Bryn Webb, MD, of our Scientific Advisory Board, to collaborate with OpenEvidence AI. The MSF's goal is to help review, edit, and accurately share information about Moebius syndrome.

MOEBIUS SYNDROME FOUNDATION REGISTRY

As part of our ongoing commitment to advancing understanding of Moebius syndrome, we would like to share updates on research surveys now available through the Moebius Syndrome Foundation Research Registry. This month, our focus is on **recurrent surveys**. One year after a Participant has entered information into a Registry survey, an update request will be received by the Participant. This prompts the Participant to enter information about new issues that have occurred in the past year. This will help us learn more about the natural progression of many aspects of Moebius syndrome, which is one of the fundamental goals of this long-term research effort. The updates for the surveys are easy and can be completed quickly in most cases.

Your engagement and participation in these research efforts can significantly shape future findings and lead to better support resources for the community.

For detailed information, visit the Moebius Syndrome Foundation Research Registry at https://www.moebius.iamrare.org. Questions? Please contact us by email at research@moebiussyndrome.org or call (844) 663-2487 ext. 705.

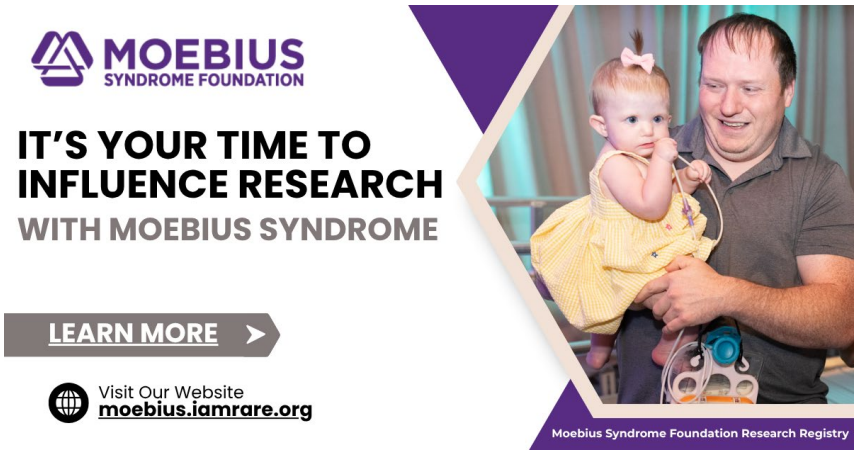
CURRENT RESEARCH PROJECTS UNDERWAY WITH GRANT FUNDING FROM MOEBIUS SYNDROME FOUNDATION

In 2025, the Moebius Syndrome Foundation awarded $100,000 grants to each of the following studies:

Learning from Adults and Parents of Children Living with Moebius Syndrome: A Rigorous Qualitative Study
Rachel Grob, Ph.D., MA (primary investigator), and Jane Evered, Ph.D., RN (co-investigator), University of Wisconsin - Madison

Leveraging Community and Professional Expertise to Accurately Characterize Early Communication Development in Children with Moebius Syndrome
Elizabeth Norton, Ph.D. (primary investigator), Judith Licht, Ph.D. candidate (co-investigator), Megan Roberts, Ph.D., CCC-SLP (co-investigator), Northwestern University

We look forward to study updates at our national conference in Pittsburgh, PA, July 22-25, 2027. Updates will also be posted to the Moebius Syndrome Foundation Research Funding page and shared in future newsletters, so our community stays informed of outcomes.



NORD®
• 2026 MEMBER •
PLATINUM

FUNDRAISERS *we love!*

AGGRESSIVELY KIND CONCERT

The Aggressively Kind hosted a successful charity concert on May 9 in Columbia, MD, bringing together friends, family, and members of the Moebius community to raise more than $1,500 for the Moebius Syndrome Foundation. Big thanks to Shelby Kennedy, volunteer Regional Coordinator for Maryland, Washington, DC, & Delaware for sharing her musical talent and organizing this event.

MULLIGANS FOR MOEBIUS

MSF Board Director Michael Mendenhall organized a very successful golf tournament fundraiser to benefit the Moebius Syndrome Foundation in Las Vegas, NV on July 25. Eighty golfers participated, and the event raised over $40,000. A special thank to Michael and his family for all their hard work to make this possible!

WE'RE FUNDRAISING WITH SEE'S CANDIES!

Starting November 1, help us raise $1,000+ for the Moebius Syndrome Foundation. Be on the lookout for a link to our virtual fundraiser. We'll share it on social media and through email.

MOVES FOR MOEBIUS

The fourth annual Moves for Moebius campaign has been taking place this September and October, and we are so "moved" by your support! People from across the country walked, rolled, and strolled to raise awareness and support for the Moebius Syndrome Foundation. Our goal is to raise $20,000 for this campaign. If you've not had an opportunity to donate to an individual, team, or just the general campaign, it's not too late. We are grateful for the outpouring of support we've seen, and hope you'll join us.

Our Colorado/Wyoming walk brought together 36 friends, family members, and supporters for an afternoon of connection, awareness, and fundraising for the Moebius Syndrome Foundation. A big thank you to volunteer Regional Coordinator Kacie Nievesmassol for organizing the event, and to everyone who came out and made it such a meaningful day for our community!

with gratitude

The mission of the Moebius Syndrome Foundation is to provide information and support to individuals with Moebius syndrome and their families, promote greater awareness and understanding of Moebius syndrome, and to advocate for scientific research to advance the diagnosis and treatment of Moebius syndrome and its associated conditions. All listed donations were made March 16, 2026 – September 15, 2026..

IN MEMORY OF

Nancy Child
Marian and Frank Setzman

Joan Cornell
Myra Enright

Jocelyn Everly Ferrante
Therese Ferrante Ferry
Matt Kuhn
Sandhya Lohithakshan
Venkatesh Manickavachagan
Robert P. Medow
Diane Morriss
David Tate
Jennifer Whitman
Amy and Gary Willow
Stephanie Winebarger

Anita Joyce Ford
Donna Hegyi-Gioia
Aileen Hower
Ivy Hill Cemetery
Robert P. Medow

Alice Profumo Goeldner
*William Goeldner**
Peter L. Profumo

Celeste Jasmyn Hayden
*Jolene Eriksen**

Mark Mudrick
Marian and Frank Setzman

Phyllis Paper
Jan Fisher

Abagail Elizabeth Simmons
Cheyenne and Joseph Nece
Jennifer Whitman

Antonius (Tony) A. Smolders
Big Blue Bubble

IN HONOR OF

Natalie Abbott
Marcia & Wayne Abbott

Aggressively Kind Concert
Elsa Arteaga
Marcia Cooper
Bob Dombrowski
Brett Dombrowski
Liz Goldstein
Stephen Kennedy
Michael Kokoski
Laura Lehman
Bill Lehrman
Brian Martin
Becki Peyton
Amy Schuster
Naseem Tarmohamed
Shelly Winston
Anonymous
Anonymous

Amélie
*Basil Nininger**

Hamilton Bachman
*Destiny Bachman**

Emily Baldridge
David Baldridge

Ben and Henry Becker
Matthew Becker

Roland Bienvenu
Dr. Anne Broussard

Lily Cicconi
Anonymous

Jackson Clark
*Alan and Dana Hyams**

Malakai Colyer
Summer Colyer

Rhys Fancher
Jeff and Barbara Hinshaw

Scotland Fernandez
Kyla Clarke

Jan Fisher
Susan Bayer
Nancy Chang
Helen Cohn
Jeannette Kaelin
Jan Klatskin
Della Levy
Barbara Marcus
Roni Ogin
Cassandra Robbins
Joyce and David Spiegler

Abigail Fulmore
Stacey Fulmore

Addison Garvey
Bounce! Family Entertainment Center

Molly Goldsmith
Carly Gershone
Steven Gershone
Ellen Kowitt
Katie Wexler

Shelby Kennedy
Maureen Connors
Lauren Disciullo
Darlene Dyer
Kim Hokanson
David Miranda

Ryan Larsen
James Larsen

Elizabeth Lawrence
Amanda and Keith Davis

Dallas Lewallen
Marilyn Donofrio
Andrea Enright
Jesse Franklin
Sandra Hanson
Kimberlee Hyp
Lindi Lewallen
Renee Peters
Ashley Roop
Bea Sloan
Carol Van Bramer
Michael Voigt
Anne Weber
Colleen Weber
Jerilyn Wright

Miriam Licht
Ronn Medow

Magical Moebius Pumpkins
Susan Scotten

Rebecca Maher
Novartis Pharmaceuticals Charity

Cheryl Manning
Give Lively Foundation

Allen Marshall
Linda Marshall

Megan Menard
Angela Coakley

Tina Michael
Jennifer Whitman

The Mob(ius) Squad
Jan Fisher
Molly Goldsmith
Robert Martin
Jennifer Whitman

Mulligans for Moebius
Melissa Mauer
Polar Shades

Khaleb Phillips
Shamika Carter
Tyrell Crease
Garnell Dickerson
Robin Fulks
Marlena Goode
Keneidra Hargrove
Ushelor Yvette Lowery
Aisha Mason
Lynette Mathis
KiOwana Phillips
Geletta Shavers
Angelia Sutton

Sloan Preisler
Samara and Seth Preisler

Kari Storm
*Tim and Laurie Argall**
Robert C. Argall Sr.

Lucas Miguel Torres
*Carlos Torres**

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Jennifer Whitman
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Erica Klauber
Robert and Chelle Medow

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Cindy Joy
Erica Klauber
Jonothan Latiak
Dallas Lewallen
*Tina Michael**
Stephen Miller
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Jared Williams
James and Brittany Yeager

ORGANIZATIONS

Brother Chimp Brewing Company
Fidelity Charitable
Give Lively Foundation
I.U.O.E.
PayPal Giving Fund
Progressive Insurance Foundation
R & L Management
Anonymous (2 gifts)

A special note of thanks goes to our Sustaining Members who have committed to making monthly contributions. They are marked with a * (star) by their names.

One Tuesday + One Community = TOGETHER TUESDAY

save the date **TUESDAY, DECEMBER 1, 2026**

There's a stretch of days at the end of November that belongs to doorbusters and shopping carts. Black Friday. Small Business Saturday. Cyber Monday. Then comes the one we look forward to all year.

Here is what you made possible this year. When a parent called us a few days after a diagnosis, concerned and unsure of where to turn for answers, someone picked up. We talked through their questions, and thanks to research funded by the Moebius Syndrome Foundation, we were able to give some real answers and point them towards resources. That is one call. Our research and community outreach initiatives have provided hope and helpful solutions to many people this year, and none of it came from a grant or a government line item. It came from this community and the people who love it.

ON TUESDAY, DECEMBER 1, WE ARE ASKING YOU TO DO IT AGAIN.

YOUR GIFT GOES TWICE AS FAR.

Our Board of Directors will once again be offering a Giving Tuesday Match Challenge. Your $100 gift becomes $200. A $250 gift becomes $500! Double your impact on Giving Tuesday! Be sure to also check with your employer for matching opportunities and help your gift go even further.

Matched gifts are limited to the challenge total, so giving early on December 1 makes sure yours is one of them. *The final match amount will be announced by November 15 in the November e-newsletter and on social media. Stay tuned!*

Thank you for every gift you have already given, and for the ones we will never know about, the conversations you started and the people you told.

WHAT YOUR MATCHED GIFT DOES:

$50 becomes $100 and provides new member packets for 5 newly diagnosed families.

$100 becomes $200 and supports community virtual meetings and gatherings for a year.

$250 becomes $500 and makes it possible to translate and produce an educational booklet into Spanish.

$500 becomes $1,000 and allows us to support 3 to 4 regional events per year.

THREE WAYS TO SHOW UP ON DECEMBER 1:

Give, and give early. Online at moebiussyndrome.org/give, or mail a check to the address below with "Giving Tuesday" in the memo line.

Moebius Syndrome Foundation
1312 17th Street, #976
Denver, CO 80202

Share your story. Post a photo and say why this matters to you. Use #GivingTuesday and #MoebiusSyndrome. Awareness travels fastest when it comes from you.

Mark your calendar. Circle December 1 now, before December does what December does. Let us know if you're planning to participate! We'd love to hear from you.

WE ARE INCREDIBLY GRATEFUL TO OUR DONORS. EACH GIFT HELPS US TO PROVIDE A BETTER FUTURE FOR INDIVIDUALS WITH MOEBIUS SYNDROME.

Scan QR code to donate online at: moebiussyndrome.org/donate/

PayPal paypal.me/moebiussyndrome

Mail a check to:
Moebius Syndrome Foundation
1312 17th Street, #976
Denver, CO 80202

Platinum Transparency 2026 Candid.

Please let us know if your gift is in honor or memory of someone.

MOEBIUS SYNDROME FOUNDATION

1312 17th Street #976
Denver, Colorado 80202

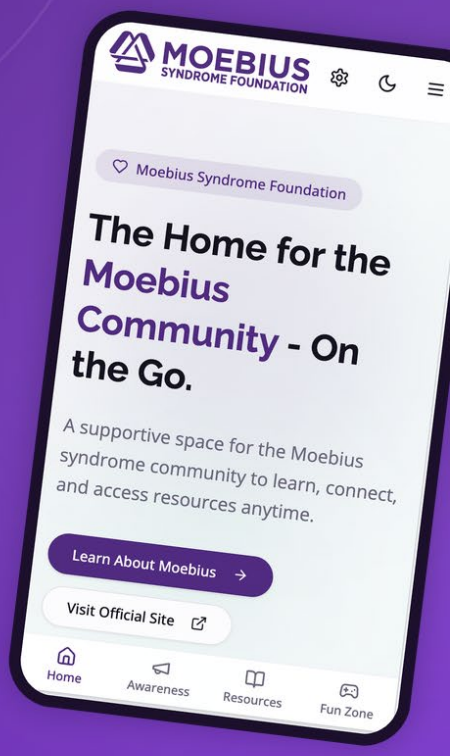
(844) 663-2487 or (844) MOEBIUS
info@moebiussyndrome.org

MOEBIUS SYNDROME FOUNDATION

NEW APP

Moebius Connect

Resources, information, and connection for the Moebius community, anytime.

Download on the App Store

GET IT ON Google Play